FDA 510(k) Premarket Notification

K810251 SYRINGE, ENT

CLEARED

Mar 11, 1981
41 days to decision

K810251 · Product code: **KYZ** · General Hospital
Source: https://www.510kdatabase.net/k810251/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Irrigating (non Dental) (KYZ)
Date received	Jan 29, 1981
Decision date	Mar 11, 1981
Days to decision	41 days
Third-party review	No

APPLICANT

Company	**Megaplast, Inc.**
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1981-1982